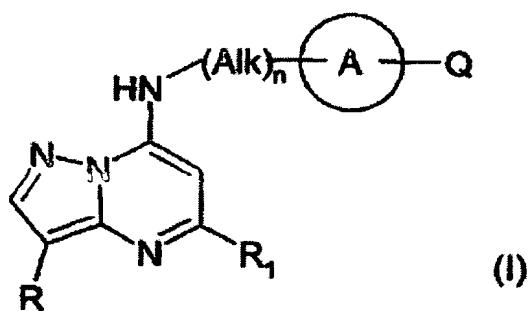


The listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Currently Amended) ~~The use of a A compound of formula (I) or a salt, N-oxide, hydrate or solvate thereof, in the preparation of a composition for inhibition of kinase activity:~~



wherein

Ring A is an optionally substituted carbocyclic or heterocyclic radical,

Alk represents an optionally substituted divalent C<sub>1</sub>-C<sub>6</sub> alkylene radical;

n is 0 or 1 ;

Q represents a radical of formula -(Alk<sup>1</sup>)<sub>p</sub>-(X)<sub>r</sub>-(Alk<sup>2</sup>)<sub>s</sub>-Z wherein in any compatible combination

Z is hydrogen or an optionally substituted carbocyclic or heterocyclic ring,

Alk<sup>1</sup> and Alk<sup>2</sup> are optionally substituted divalent C<sub>1</sub>-C<sub>6</sub> alkylene radicals which may contain a -O-, -S- or -NR<sup>A</sup>-link, wherein R<sup>A</sup> is hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl,

X represents -O-, -S-, -(C=O)-, -(C=S)-, -SO<sub>2</sub>-, -SO-, -C(=O)O-, -OC(=O)-, -C(=O)NR<sup>A</sup>-, .NR<sup>A</sup>C(=O)-, -C(=S)NR<sup>A</sup>-, -NR<sup>A</sup>C(=S)-, -SO<sub>2</sub>NR<sup>A</sup>-, -NR<sup>A</sup>S0<sub>2</sub>-, -OC(=O)NR<sup>A</sup>-, -NR<sup>A</sup>C(=O)O-, or -NR<sup>A</sup>- wherein R<sup>A</sup> is hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl, and

p, r and s are independently 0 or 1,

R<sub>1</sub> represents a radical -(Alk<sup>3</sup>)<sub>a</sub>-(Y)<sub>b</sub>-(Alk<sup>4</sup>)<sub>d</sub>-B wherein a, b and d are independently 0 or 1,

Alk<sup>3</sup> and Alk<sup>4</sup> are optionally substituted divalent C<sub>1</sub>-C<sub>3</sub> alkylene radicals,

Y represents a monocyclic divalent carbocyclic or heterocyclic radical having from 5 to 8 ring atoms, -O-, -S-, or-NR<sup>A</sup>- wherein R<sup>A</sup> is hydrogen or C<sub>1</sub>-C<sub>6</sub> alkyl,

B represents hydrogen or halo, or an optionally substituted monocyclic carbocyclic or heterocyclic ring having from 5 to 8 ring atoms, or in the case where Y is -NR<sup>A</sup>- and b is 1, then R<sup>A</sup> and the radical-(Alk<sup>4</sup>)<sub>d</sub>-B taken together with the nitrogen to which they are attached may form an optionally substituted heterocyclic ring,

R represents hydrogen, halo, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>1</sub>-C<sub>6</sub> alkoxy, C<sub>1</sub>-C<sub>6</sub> alkylthio, phenyl, benzyl, cycloalkyl with 3 to 6 ring atoms, or a monocyclic heterocyclic group having 5 or 6 ring atoms.

2. (Currently Amended) The use compound as claimed in claim 1 wherein ring A is an optionally substituted monocyclic aryl or heteroaryl radical.

3. (Currently Amended) The use compound as claimed in claim 2 wherein ring A is phenyl, naphthyl, 2-, 3- and 4-pyridyl, 5-pyrimidinyl, 2- and 3-thienyl, 2- and 3-furyl, piperazinyl, pyrrolidinyl, or thiazolinyl.

4. (Currently Amended) The ~~use compound~~ as claimed in claim 1 wherein ring A is phenyl.
5. (Currently Amended) The ~~use compound~~ as claimed in ~~any of the preceding claims~~ claim 1 wherein ring A is unsubstituted or substituted by methyl, ethyl, methylenedioxy, ethylenedioxy, methoxy, ethoxy, methylthio, ethylthio, hydroxy, hydroxymethyl, hydroxyethyl, mercapto, mercaptomethyl, mercaptoethyl, amino, mono- or di-methylamino, mono- or di-ethylamino, fluoro, chloro, bromo, cyano, N-morpholino, N-piperidinyl, or N-piperazinyl, the latter being optionally C<sub>1</sub>-C<sub>6</sub> alkyl- or benzyl-substituted on the free ring nitrogen, dimethylaminosulfonyl, phenylsulfonyl or phenoxy.
6. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 3~~ claim 1 wherein Q is hydrogen and the ring A is 4-(dimethylaminosulfonyl)-phenyl, 4-(phenylsulfonyl)-phenyl, 4-(phenoxy)-phenyl, 3-chloro-4-(dimethylaminosulfonyl)-phenyl, 3-chloro-4(phenylsulfonyl)-phenyl, 3-chloro-4-(phenoxy)-phenyl, 3-methoxy-4(dimethylaminosulfonyl)-phenyl, 3-methoxy-4-(phenylsulfonyl)-phenyl, or 3-methoxy-4-(phenoxy)-phenyl.
7. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1 wherein n is 1 and Alk is CH<sub>2</sub>-, -CH<sub>2</sub>CH<sub>2</sub>-, -CH<sub>2</sub>CH(CH<sub>3</sub>)-, -CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>-, -CH=CH-, -CH<sub>2</sub>CH=CH-, -CH<sub>2</sub>CH=CHCH<sub>2</sub>-, -CH=CHCH=CH-, -C=C-, -CH<sub>2</sub>C=C-, or -CH<sub>2</sub>C=CCH<sub>2</sub>-.
8. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1 wherein n is 1 and Alk is -CH<sub>2</sub>-.
9. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1 wherein n is 0.
10. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1 wherein each of p, r and s is 0, and Z is hydrogen.
11. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1

wherein p, r and s are each 0, and Z is an optionally substituted monocyclic carbocyclic or heterocyclic ring.

12. (Currently Amended) The ~~use compound~~ as claimed in claim 11 wherein Z is an optionally substituted phenyl, cyclopentyl, cyclohexyl, pyridyl, morpholino, piperidinyl, or piperazyl ring.

13. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 5~~ claim 1 wherein one or more of p, r and s is 1, and Z is hydrogen or an optionally substituted monocyclic carbocyclic or heterocyclic ring.

14. (Currently Amended) The ~~use compound~~ as claimed in claim 13 wherein p, and/or s, or both are each 1 and r is 0.

15. (Currently Amended) The ~~use compound~~ as claimed in claim 13 wherein each of p, r, and s is 1.

16. (Currently Amended) The ~~use compound~~ as claimed in claim 13 wherein p and s are each 0 and r is 1.

17. (Currently Amended) The ~~use compound~~ as claimed in claim 16 wherein X is -SO<sub>2</sub>-, -O-, a sulfonamide radical -NR<sup>A</sup>SO<sub>2</sub>- or a carboxamide radical -NR<sup>A</sup>C(=O)- with the N atom linked to the ring A.

18. (Currently Amended) The ~~use compound~~ as claimed in claim 13 wherein p is 0, r is 1, s is 1 or 0, and X is a sulfonamide radical -NR<sup>A</sup>SO<sub>2</sub>- or a carboxamide radical -NR<sup>A</sup>C(=O)- with the N atom linked to the ring A.

19 (Currently Amended) The ~~use compound~~ as claimed in claim 17 ~~or claim 18~~ wherein R<sup>A</sup> is hydrogen or methyl.

20. (Currently Amended) The ~~use compound~~ as claimed in claim 18 or ~~claim 19~~ wherein s is 1 and Z is hydrogen.
21. (Currently Amended) The ~~use compound~~ as claimed in claim 18 or ~~claim 19~~ wherein s is 0 and Z is an optionally substituted monocyclic carbocyclic or heterocyclic ring.
22. (Currently Amended) The ~~use compound~~ as claimed in claim 21 wherein Z is optionally substituted phenyl.
23. (Currently Amended) The ~~use compound~~ as claimed in ~~any of the preceding claims~~ ~~claim 1~~ wherein in the radical R<sub>1</sub> a, b and d are all 0.
24. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 22~~ ~~claim 1~~ wherein in the radical R<sub>1</sub> a and d are each 0 and b is 1.
25. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 1 to 22~~ ~~claim 1~~ wherein in the radical R<sub>1</sub> b is 0 and at least one of a and d is 1.
26. (Currently Amended) The ~~use compound~~ as claimed in ~~any of claims 23 to 25~~ ~~claim 23~~ wherein in the radical R<sub>1</sub>, B is an optionally substituted monocyclic carbocyclic or heterocyclic ring.
27. (Currently Amended) The ~~use compound~~ as claimed in claim 26 wherein B is an optionally substituted cyclopentyl, cyclohexyl, phenyl, 2-,3-, or 4-pyridyl, 2-, or 3-thienyl, 2-, or 3-furanyl, pyrrolyl, pyranyl, or piperidinyl ring.
28. (Currently Amended) The ~~use compound~~ as claimed in claim 27 wherein optional substituents are selected from methyl, ethyl, methoxy, ethoxy, methylenedioxy, ethylenedioxy, methylthio, ethylthio, hydroxy, hydroxymethyl, hydroxyethyl, mercapto, mercaptomethyl, mercaptoethyl, amino, mono- and di-methylamino, mono- and diethylamino, fluoro, chloro, bromo, cyano, N-morpholino, N-piperidinyl, N-piperazinyl.

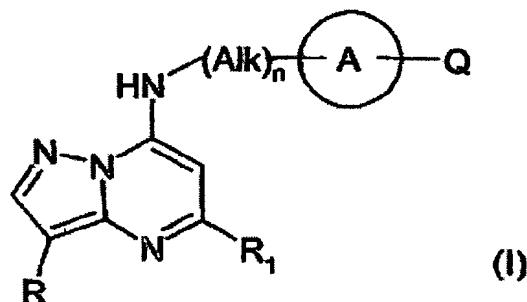
29. (Currently Amended) The ~~use compound as claimed in any of claims 1 to 22~~ claim 1 wherein R<sub>1</sub> is optionally substituted cyclohexyloxy; cyclohexylamino; cyclohexylmethyl, or piperidin-1-ylmethyl.
30. (Currently Amended) The ~~use compound as claimed in any of claims 1 to 22~~ claim 1 wherein R<sub>1</sub> is 4-aminocyclohexyloxy; 4-aminocyclohexylamino; 4-hydroxycyclohexylamino, 4-aminocyclohexylmethyl, or 4-aminopiperidin-1-ylmethyl.
31. (Currently Amended) The ~~use compound as claimed in any of the preceding claims~~ claim 1 wherein R is hydrogen, chloro, bromo methyl, ethyl, n-propyl, iso-propyl, n-, sec- or tert-butyl, methoxy, methylthio, ethoxy, ethylthio, or a phenyl, benzyl, cyclopropyl, cyclopentyl, cyclohexyl, 2-, 3-, or 4- pyridyl, phenyl, pyridyl, morpholino, piperidinyl, or piperazyl ring.
32. (Currently Amended) The ~~use compound as claimed in any of claims 1 to 30~~ claim 1 wherein R is chloro, bromo, cyclopentyl, cyclopropyl or isopropyl.
33. (Currently Amended) The ~~use compound as claimed in claim 1~~ wherein in the compound of formula (I) n is 0, ring A is optionally substituted phenyl, Q is dimethylaminosulfonyl, phenylsulfonyl or phenoxy; R<sup>1</sup> is 4-aminocyclohexyloxy, 4aminocyclohexylamino, 4-hydroxycyclohexylamino, 4-aminocyclohexylmethyl, or 4-aminopiperidin-1-ylmethyl, and R is chloro, bromo, cyclopentyl, cyclopropyl or isopropyl.
34. (Currently Amended) A method of treatment of diseases or conditions mediated by excessive or inappropriate kinase activity in mammals, ~~particularly humans, which method comprises comprising~~ administering to the mammal an amount of a compound of formula (I) as defined in ~~any of the preceding claims~~ claim 1, or a salt, hydrate or solvate thereof, effective to inhibit said kinase activity.
35. (Canceled)
36. (Currently Amended) The ~~use as claimed in any of claims 1 to 33~~, a method as claimed in

claim 34, or a compound for use as claimed in claim 35, wherein the kinase activity is CDK2 activity, and/or PDK1 activity, and/or CHK1 activity, or combinations thereof.

37. (Currently Amended) The use as claimed in any of claims 1 to 33, a method of treatment as claimed in claim 34, or a compound for use as claimed in claim 35 wherein the kinase activity is associated with cancer, psoriasis or restenosis.

38. (Currently Amended) A pharmaceutical composition comprising a compound of formula (I) as defined in any of claims 1 to 32 claim 1, or a salt, N-oxide, hydrate or solvate thereof, together with a pharmaceutically acceptable carrier.

39. (Currently Amended) A compound of formula (I), or a salt, N-oxide, hydrate or solvate thereof,



wherein n is 0, ring A is optionally substituted phenyl, Q is dimethylaminosulfonyl, phenylsulfonyl or phenoxy, R<sup>1</sup> is 4-aminocyclohexyloxy; 4-aminocyclohexylamino; 4-hydroxycyclohexylamino; 4-aminocyclohexylmethyl, or 4-aminopiperidin-1-ylmethyl, and R is chloro, bromo, cyclopentyl, cyclopropyl or isopropyl.

40 (Currently Amended) A pharmaceutical composition as claimed in claim 38 or claim 39 together with a pharmaceutically acceptable carrier.